PTO-1449 REPRODUCED

INFORMATION DISCLOSURE CITATION IN AN APPLICATION

May 23, 2001

(Use several sheets if necessary)

ATTORNEY DOCKET NO.	APPLICATION NO.
3028.1000-000	09/590,211
APPLICANT	
Guy A. Rouleau and Bernard Brais	
FILING DATE	**GROUP**
June 8, 2000	1632

U.S. PATENT DOCUMENTS

EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUB-CLASS	FILING DATE IF APPROPRIATE
	AA						
	AB						

FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB-CLASS	TRANSLATION YES	TRANSLATION NO
✓	AL	WO98/31800	23 JUL 98	PCT				
✓	AM	WO99/29896	17 JUN 99	PCT				
	AN							
	AO							

OTHER DOCUMENTS ***(Including Author, Title, Date, Pertinent Pages, Etc.)***

✓	AR ✓	Akarsu, A.N., et al., "Genomic Structure of HOXD13 Gene: A Nine Polyalanine Duplication Causes Synpolydactyly in Two Unrelated Families," *Human Molecular Genetics*, 5(7): 945-952 (1996).
	AS ✓	Bienroth, S, et al., "Assembly of a Processive Messenger RNA Polyadenylation Complex," *The EMBO Journal*, 12(2): 585-594 (1993).
	AT ✓	Brais, B., et al., "Using the Full Power of Linkage Analysis in 11 French Canadian Families to Fine Map the Oculopharyngeal Muscular Dystrophy Gene," *Neuromuscular Disorder* 7(1):S70-S74 (1997).
	AU ✓	Brais, B, et al., "The Oculopharyngeal Muscular Dystrophy Locus Maps to the Region of the Cardiac α and β Myosin Heavy Chain Genes on Chromosome 14q11.2-q13," *Human Molecular Genetics*, 4(3): 429-434 (1995).
	AV ✓	Davies, S.W., "Formation of Neuronal Intranuclear Inclusions Underlies the Neurological Dysfunction in Mice Transgenic for the HD Mutation," *Cell*, 90:537-548 (1997).
	AW ✓	DiFiglia, M, et al., "Aggregation of Huntingtin in Neuronal Intranuclear Inclusions and Dystrophic Neurites in Brain," *Science*, 277: 1990-1993 (1997).
	AX ✓	Evans, G.A, et al., "High Efficiency Vectors for Cosmid Microcloning and Genomic Analysis," *Gene*, 79:9-20 (1989).
	AY ✓	Forood, B., et al., "Formation of an Extremely Stable Polyalanine β-Sheet Macromolecule," *Biochem. And Biophysical Res. Communications*, 211(1): 7-13 (1995).
	AZ ✓	Krause, S., et al., "Immunodetection of Poly(A) Binding Protein II in the Cell Nucleus," *Experimental Cell Res.*, 214: 75-82 (1994).

EXAMINER	DATE CONSIDERED
Joe Woitach	7/8/04

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OTHER DOCUMENTS ***(Including Author, Title, Date, Pertinent Pages, Etc.)***

AR2	Mundlos, S., et al., "Mutations Involving the Transcription Factor CBFA1 Cause Cleidocranial Dysplasia," *Cell*, 89: 773-779 (1997).
AS2	Editorials, "DNA-Triplet Repeats and Neurologic Disease," *The New England Journal of Med.*, 335(16): 1222-1224 (1996).
AT2	Scherzinger, E., et al, "Huntingtin-Encoded Polyglutamine Expansions Form Amyloid-Like Protein Aggregates In Vitro and In Vivo," *Cell* 90: 549-558 (1997).
AU2	Tome, M.S., et al., "Nuclear Inclusions in Oculopharyngeal Dystrophy," *Act Neuropathol.* 49: 85-87 (1980).
AV2	Muragaki, Y., et al., "Polyalanine Expansion in Synpolydactyly Might Result from Unequal Crossing-Over of HOXD13," *Science* 275: 406
AW2	Wells, R.D., "Molecular Basis of Genetic Instability of Triplet Repeats," *The Journal of Biological Chem.* 271(6): 2875-2878 (1996).
AX2	Wahle, E., et al., "Mammalian Poly(A)-Binding Protein II," *J. of Biological Chem.*, 268(4): 2937-2945 (1993).
AY2	Wahle E., "A Novel Poly(A)-Binding Protein Acts As a Specificity Factor in the Second Phase of Messenger RNA Polyadenylation," *Cell*, 66: 759-768 (1991).
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AS3	Brais, B., et al., "Short GCG Expansions in the PABP2 Gene Cause Oculopharyngeal Muscular Dystrophy," *Nature Genetics* 18: 164-167 (1998).

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JL	AT3 ✓	Lamartine, J. et al., "Cloning Sequencing and Chromosomal Assignment of a New cDNA Clone to Xq12-q13 and 14q11," EMBL Data Base Accession Number U12206 (1995).
	AU3 ✓	"National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index," EMBL Data Base Accession Number AA618589 (1997).
	AV3 ✓	Sullivan, T.B.T., et al., "Oculopharyngeal Muscular Dystrophy (OPMD) - Report and Genetic Studies of an Australian Kindred," *Clinical Genetics*, 51: 52-55 (1997).
	AW3 ✓	Bouchard, J.P. *et al.*, "A Simple Test for the Detection of Dysphagia in Members of Families with Oculopharyngeal Muscular Dystrophy (OPMID)," Can.J. Neurol. Sci. 19(2):296-297 (1992).

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